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In the Claims:

Please amend the claims by replacing all prior versions of the claims pursuant to 37 C.F.R. §1.121 as modified by 68 Fed. Reg. 38611 (June 30, 2003) as follows:

- 1-20. (Previously Canceled)
- 21. (Previously Presented) The method of claim 50, wherein F_1 corresponds to a segment of amino acid residues from within N-terminal residues 1-10 of SCF (SEQ ID NO:1), F_2 corresponds to a segment of amino acid residues from within residues 79-95 of SCF, and F_3 corresponds to a segment of amino acid residues located within three amino acid residues of amino acid residue 127 of SCF, and where, in X_n , X_m , and X_p respectively, n=0-5, m=0-5 and p=3-8 amino acid residues.
- 22. (Previously Presented) The method of claim 50, wherein F_1 , F_2 , and F_3 have been selected by bacterial phage display for optimal receptor binding.
- 23-25. (Previously Canceled)
- 26. (Previously Presented) The method of claim 50, wherein the organic polymer is polyethyleneglycol (PEG) comprising the structure $H[OCH_2CH_2]_nOH$, wherein n is 10-20.
- 27. (Previously Presented) The method of claim 50, wherein the capping moiety is a thiol-reactive group.
- 28-47. (Previously Canceled)

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- 48. (Currently Amended) A method for designing a compound capable of binding to a Stem Cell Factor-binding site of a Kit receptor Stem Cell Factor receptor comprising the steps of:
 - a) determining the 3-D structure of a fragment of a Stem Cell Factor (SCF) by computing atomic coordinates from X-ray diffraction data of a crystal of the fragment of SCF, wherein the fragment of SCF consists of consecutive amino acids the sequence of which is set forth in SEQ ID NO:1;
 - b) determining a Kit receptor identifying a Stem Cell Factor receptor-binding site on the fragment of SCF based on the 3-D structure of the SCF fragment; and
 - c) and designing a compound capable of binding to the Stem Cell Factor-binding site of the Stem Cell Factor receptor of the Kit receptor based on a 3-D structure shape complementarity or estimated interaction energy of the Stem Cell Factor receptor-binding site on the fragment of SCF.
- 49. (Previously Canceled)
- 50. (Currently Amended) The method of claim 48, wherein the designed compound capable of binding to a Kit Stem Cell Factor receptor comprises two ligand heads linked by a linker molecule, wherein the linker molecule is an organic polymer attached at each end to a separate capping moiety, each capping moiety attached in turn to a single ligand

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head via a cysteine residue, wherein the ligand head comprises the elements F_1 - X_n - F_L (Cys)- X_m - F_2 - X_p - F_3 , wherein F_1 , F_2 and F_3 are peptides each comprising amino acid sequences corresponding to consecutive amino acid residues of SCF (SEQ ID NO:1), X_n and X_p are peptides of n and p amino acid residues respectively, F_L is the cysteine residue and each element is linked to the next via a peptide bond.

51. (Previously Presented) The method of claim 27, wherein the thiol-reactive group is N-ethyl malemide.